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A Panchromatic, Near Infrared Ir(III) Emitter Bearing a Tripodal C^N^C ligand as a Dye for Dye-Sensitized Solar Cells

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Abstract

The synthesis of a new complex of the form [Ir(C^N^C)(N^N)Cl] [where C^N^C = 2-(bis(4-(*tert*-butyl)phenyl)methyl)pyridinato (d*t*Bubnpy, **L1**) and N^N is diethyl [2,2'-bipyridine]-4,4'-dicarboxylate (deeb)] is reported. The crystal structure reveals an unusual tripodal tridentate C^N^C ligand forming three six-membered rings around the iridium center. The photophysical and electrochemical properties suggest the use of this complex as a dye in dye-sensitized solar

cells. Time-Dependent Density Functional Theory (TD-DFT) calculations have been used to reveal the nature of the excited-states.

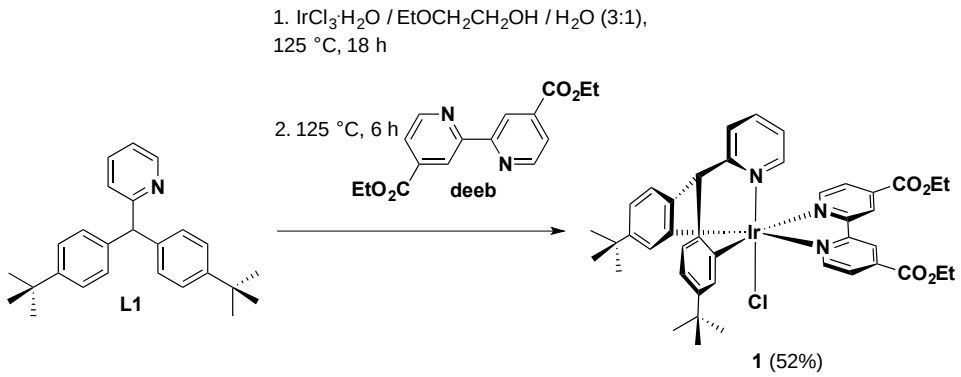
Introduction

Dye-sensitized solar cells (DSSCs) represent a promising solar cell technology. The majority of champion DSSCs, those showing power conversion efficiencies (PCE) greater than 10%, are based on ruthenium(II) complexes. Iridium(III) complexes, dominant as emitters in electroluminescent devices,[1,2] have to date fared poorly as dyes in DSSCs.[3-13] This is mainly because most iridium(III) complexes are not panchromatic, having absorption spectra that tail off by 550 nm. This induces low short circuit currents in the DSSC and as a consequence poor PCE; typically less than 4%. Indeed, there are very few examples of iridium(III) complexes with significant absorption bands going up to the red or NIR parts of the visible spectrum.[14-20,12]

We recently reported the development of tripodal C^N^C ligands, 2-benzhydrylpyridine and its derivatives, which can coordinate to iridium, forming three six-membered chelate rings through a double C-H bond activation.[21] When combined with a bidentate diimine ligand such as 4,4'-di*tert*butyl-2,2'-bipyridine (d*t*Bubpy), a family of orange-to-red emitting neutral [Ir(C^N^C)(d*t*Bubpy)Cl] complexes was formed with absorption bands tailing off at 600 nm. Herein, we report an analogous complex showing panchromatic absorption, employing an electron-poor ancillary ligand diethyl [2,2'-bipyridine]-4,4'-dicarboxylate (deeb), and study its use as a DSSC dye.

Results and Discussion

Synthesis



Scheme **1**: Scheme for the one-pot synthesis of complex **1**.

Compound **L1**[21] and deeb[22] were prepared by literature methods. Complex **1** was obtained as a black solid in 52% yield using a two-step-one-pot protocol wherein a mixture of **L1** and $IrCl_3.6H_2O$ in 2-ethoxyethanol/H_2O (3:1) was heated at reflux for 19 h followed by the addition of deeb and a further reaction time of 6 h (Scheme **1**). Complex **1** was characterized by ^{1}H and ^{13}C NMR spectroscopy, HR-ESI mass spectrometry, elemental analysis and melting point determination [see Figures **S1-6** in the Supporting Information (SI) for NMR and HR-ESI mass spectra].

Crystal Structures

Single crystals of sufficient quality of **1** were grown from CH_2Cl_2/Et_2O at -18°C, and the structure of **1** was determined by single-crystal X-ray diffraction (Figure **1,** Table **S1**). Complex **1**, [Ir(**L1**)(deeb)Cl], lies in a mirror plane; the pyridyl ring of **L1**, the iridium(III) and the chloride all lying directly in the plane. The tridentate **L1** shows a tripodal chelation motif, analogous to that seen previously.[21] The remaining coordination sphere of **1** consists of the deeb N^N ligand and a chloride anion. The arrangement of ligands is unusual, as the chloride coordinates *trans* to the

pyridine of **L1** and not *trans* to a cyclometalated carbon ligand as generally observed in tridentate complexes,[23-27] although this coordination arrangement was found in our previous complexes based on **L1**.[21] The Ir-Cl bond is 2.346(3) Å, which is within the range of distances we have previously reported for [Ir(C^N^C)(d*t*Bubpy)Cl] complexes (where C^N^C is a substituted 2-benzhydrylpyridine),[21] and is similar to the Ir-Cl distance seen in the related complex $[Ir(tpy)(dmbpy)Cl]^{2+}$ (where tpy = 2,2':6',2''-terpyridine and dmbpy = 4,4'-dimethyl-2,2'-bipyridine).[28] The $Ir\text{-}N_{N^N}$ bonds [2.134(6) Å] are in the same range as both our related [Ir(C^N^C)(d*t*Bubpy)Cl] complexes,[21] as well as the deeb complex $[Ir(topy)_2(deeb)]PF_6$ [where topyH = 2-(*p*-tolyl)pyridine].[29] The Ir-N distance involving the nitrogen from **L1** [2.038(8) Å] is markedly shorter than the $Ir\text{-}N_{N^N}$ distance. The $Ir\text{-}C_{C^N^C}$ bonds [2.027(7) Å] are shorter again than the Ir-N bonds, in agreement with our previous observations.[21] The bite angle of the N^N ligand is 76.1(3)°, in line with other iridium complexes possessing bidentate diimine ancillary ligands.[30-33,21] The C-Ir-C angle [85.2(4)°] and N-Ir-C angle within **L1** [88.0(2)°] are significantly larger due to the formation of three 6 membered chelate rings in the C^N^C ligand.

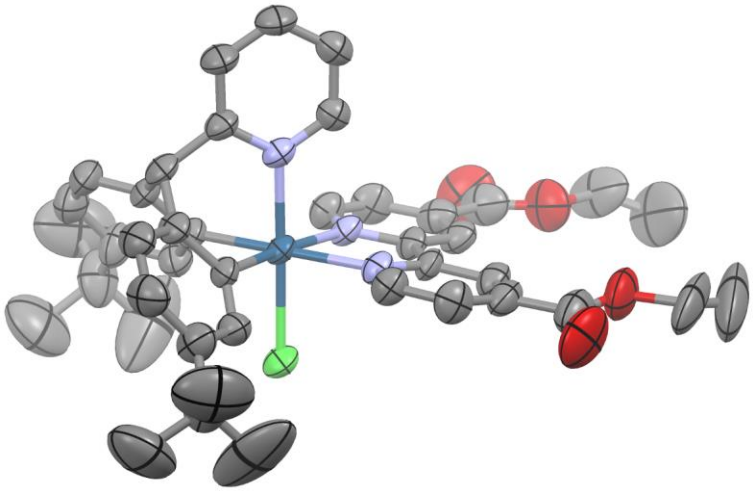


Figure **1**. Solid-state structure of complex **1**, thermal ellipsoids are drawn at the 50 % probability level. Hydrogen atoms and solvent molecules are omitted for clarity (color code: C = grey, N = purple, O = red, Cl = green and Ir = blue).

Electrochemical properties

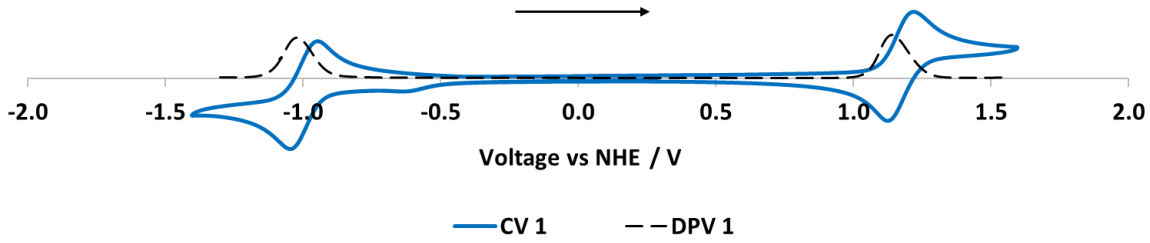


Figure **2**. Cyclic voltammograms (in blue solid lines) and differential pulse voltammetry (in dotted black lines) carried out in degassed CH_2Cl_2 at a scan rate of 100 mV s^{-1}, with Fc/Fc^+ as the internal reference, referenced to NHE (0.70 V vs. NHE).[34]

The electrochemical properties of **1** were evaluated by cyclic voltammetry (CV) and differential pulse voltammetry (DPV) in deaerated CH_2Cl_2 solution at 298 K at a scan rate of 100 mV s^{-1} using Fc/Fc^+ as the internal reference and referenced with respect to NHE (0.70 V vs. NHE).[34] The electrochemical data are summarized in Table **1** and the voltammograms are shown in Figure **2**. Complex **1** exhibits a quasi-reversible single electron oxidation wave at 1.21 V, which is assigned to the Ir(III)/Ir(IV) redox couple, with contributions from the two phenyl rings of **L1** and the chloro ligand. Compared to [Ir(**L1**)(d*t*Bubpy)Cl], **R1**, (Figure **3**, $E_{1/2}^{ox.}$ 1.04 V vs. NHE)[21] the oxidation potential in **1** is significantly anodically shifted by 170 mV, reflecting the electron-withdrawing capacity of the ethyl ester groups of the N^N ligand, which modifies the

electron density on iridium. However, the oxidation potential of **1** is less positive than that of $[Ir(ppy)_2(deeb)]PF_6$, **R2**, ($E_{1/2}^{ox.}$ = 1.57 V in deaerated MeCN vs NHE, where ppy is 2-phenylpyridinato).[35] Upon scanning to negative potential, **1** shows a single quasi-reversible reduction wave at -0.94 V, which is monoelectronic as inferred from the DPV. The electron-withdrawing effect of the ethyl ester groups of the N^N ligand results in a large anodic shift of 610 mV in the reduction wave of **1** compared to **R1** ($E_{1/2}^{red.}$ -1.58 V vs NHE).[21] Complex **R2** showed two reversible reduction waves in MeCN. The first reduction located at -0.76 V is assigned to the reduction of the deeb ligand while the second one at -1.30 V is due to the reduction of the phenylpyridinato.[35] Thus, the reduction of the deeb ligand in **1** is shifted to more negative potentials compared to the same reduction in **R2**. DFT calculations of the previously reported **R1** indicated that both the HOMO and HOMO-1 are close in energy and involve the iridium and chlorine atoms and the two phenyl rings of **L1**.[21] As can be seen in Figure **4** the same electron density distribution is found in **1**. DFT calculations also show that the three lowest unoccupied orbitals are exclusively localized on the deeb ligand in **1** (Figure **4**), while the LUMO+1 is primarily on the pyridyl of **L1** in **R1**, illustrating the stronger accepting character of deeb. The $\Delta E_{\mathrm{redox}}$ for **1** (2.18 eV) is markedly smaller than that of **R2** ($\Delta E_{\mathrm{redox}}$ = 2.33 V).[35]

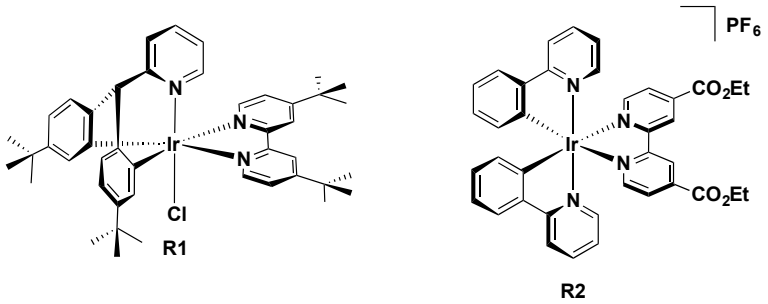


Figure **3**. Structures of reference complexes **R1** and **R2**.

Table **1**: Selected electrochemical properties of complex **1**

	Electrochemistry[a]						
	$E_{1/2}^{ox}$ / V	ΔE_p / mV	$E_{1/2}^{red}$ / V	ΔE_p / mV	ΔE_{redox}[b] / V	E_{HOMO}[c] /eV	E_{LUMO}[c] / eV
1	1.21	88	-0.97	99	2.18	-5.31	-3.13

[a] in degassed CH_2Cl_2 at a scan rate of 100 mV s^{-1} with Fc/Fc^+ as internal reference, and referenced with respect to NHE (Fc/Fc^+ = 0.70 V in CH_2Cl_2); [36,34,37] [b] ΔE_{redox} is the difference (V) between first oxidation and first reduction potentials; [c] $E_{HOMO/LUMO}$ = -[$E^{ox/red}$ vs Fc/Fc^+ + 4.8] eV.[34]

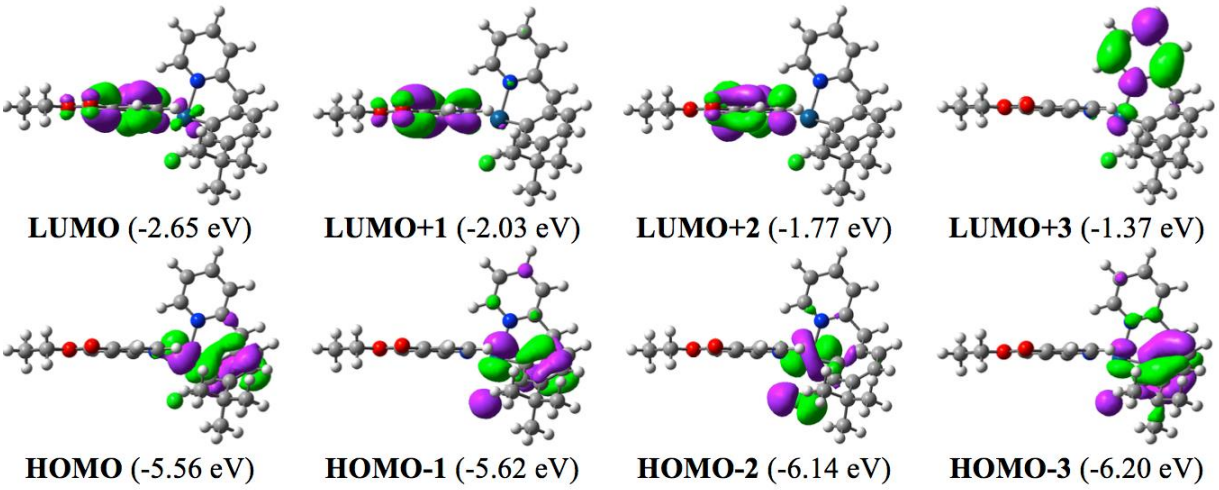


Figure **4**. Frontier molecular orbitals of **1** computed through DFT (M06 functional, see the SI for details) and represented using a contour threshold of 0.03 au.

Photophysical properties

The photophysical data for **1** recorded in CH_2Cl_2 at 298 K are shown in Figure **5** and the data summarized in Table **2**. The absorption profile of **1** differs significantly from that of **R1**. Complex **1** shows intense high-energy absorption bands (ε on the order of 3.5 × 10^4 M^{-1} cm^{-1}) below 250 nm, which are ascribed to $^1\pi$–π^* ligand-centered (^{1}LC) transitions localized on the deeb ligand. A moderately intense band (ε on the order of 1.5 × 10^4 M^{-1} cm^{-1}) at 319 nm is assigned to

a ligand-centered (LC) transition on the deeb with a small CT character (see below). Weaker bands (ε on the order of 5-6 × 10^3 and 2 × 10^3 M^{-1} cm^{-1}) in the region of 380 – 440 nm and tailing to 500 - 600 nm are attributed to a mixture of (^{1}MLCT/^{1}LLCT) and spin-forbidden (^{3}MLCT/^{3}LLCT) transitions involving the deeb ligand. Iridium(III) complexes often do not show absorption onsets lower in energy than 550 nm;[38-40] though, there are known examples of neutral Ir(III) complexes showing absorption bands beyond 550 nm.[41,25,42,43]

The assignments for complex **1** were confirmed by TD-DFT calculations (see the ESI for technical details). The two lowest singlet states, computed at 623 and 611 nm, present relatively small intensities (oscillator strengths, *f*, of 0.010 and 0.056, respectively) and mainly correspond to HOMO-1 to LUMO and HOMO to LUMO transitions. As can be seen in Figure **4**, this clearly corresponds to a mixed CT process from the metal and the phenyl rings of the C^N^C ligand towards the deeb. The following significant vertical absorption are predicted by TD-DFT at 496 nm (*f*=0.071), 456 nm (*f*=0.027) and 443 nm (*f*=0.084) and these bands can be ascribed to HOMO-2 to LUMO, HOMO to LUMO+1 and HOMO-1 to LUMO+1 transitions, respectively, and therefore all involve strong CT character towards the deeb moiety. The more intense and resolved band at 319 nm experimentally (see Table **2**) is computed at 315 nm by TD-DFT (*f*=0.162) and corresponds to a more LC excitation from a low-lying orbital centered on the deeb (and partly on chlorine atom) towards the LUMO centered on the deeb as well.

Upon photoexcitation at 420 nm, **1** exhibits a broad featureless profile, indicative of an emission with mixed CT character, with a maximum at λ_{em} = 731 nm, an emission that is significantly redshifted (99 nm, 2194 cm^{-1}) compared to **R1** (λ_{em} = 630 nm).[21] The red-shifted

luminescence is due to the presence of the presence of the π-accepting deeb. The emission of **1** is likewise red-shifted (51 nm, 2194 cm^{-1}) compared to that of **R2** (λ_{em} = 680 nm).[35] The DFT calculations returns an emission of the T_1 state at 762 nm, close to the experimental value, confirming emission from the lowest triplet excited state. The topology of this state, in terms of localization of the excess α electrons, is displayed in Figure **6**. As can be seen, the spin density is mostly localized on the Ir and Cl atoms and on the ancillary ligand, the tridentate ligand playing only a minor role in this state. This localization is consistent with the observed red-shift in emission compared to **R1** and **R2**. The measured photoluminescence quantum yield (Φ_{PL}) of **1** is 0.5%, lower than those of **R1** (6%) and **R2** (5%), a logical consequence of the energy gap law. Among near-infrared emissive cationic Ir(III) emitters with λ_{em} beyond 700 nm bearing diimines as ancillary ligand, most examples exhibit Φ_{PL} values less than 4%.[40,44-47] However, NIR-emitting neutral Ir(III) complexes of the form $[Ir(C^N)_2(O^O)]$ (where O^O a substituted β-diketonate ancillary ligand) employing highly conjugated C^N ligands have reached Φ_{PL} of up to 16%.[48,41,49] Complex **1** exhibits a multiexponential emission decay, a reflection of the large non-radiative decay rate constant.

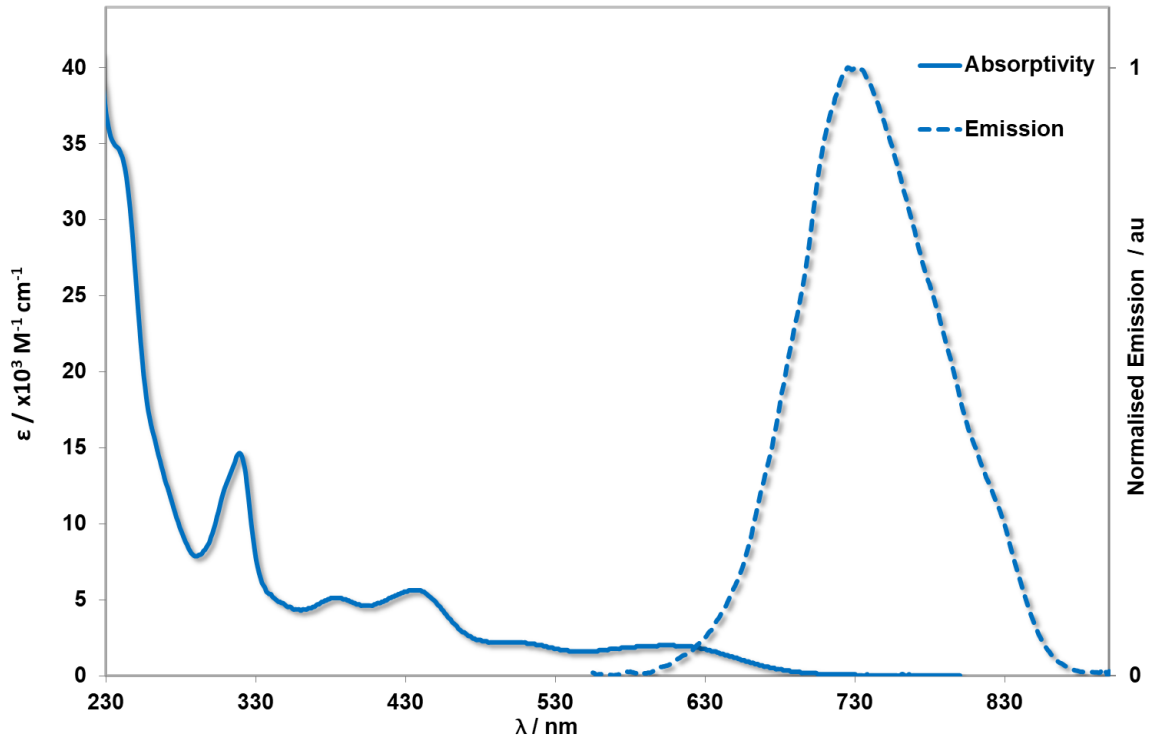


Figure **5.** The absorptivity (solid line) and photoluminescence spectra (dotted line) of **1** in CH_2Cl_2 at 298 K.

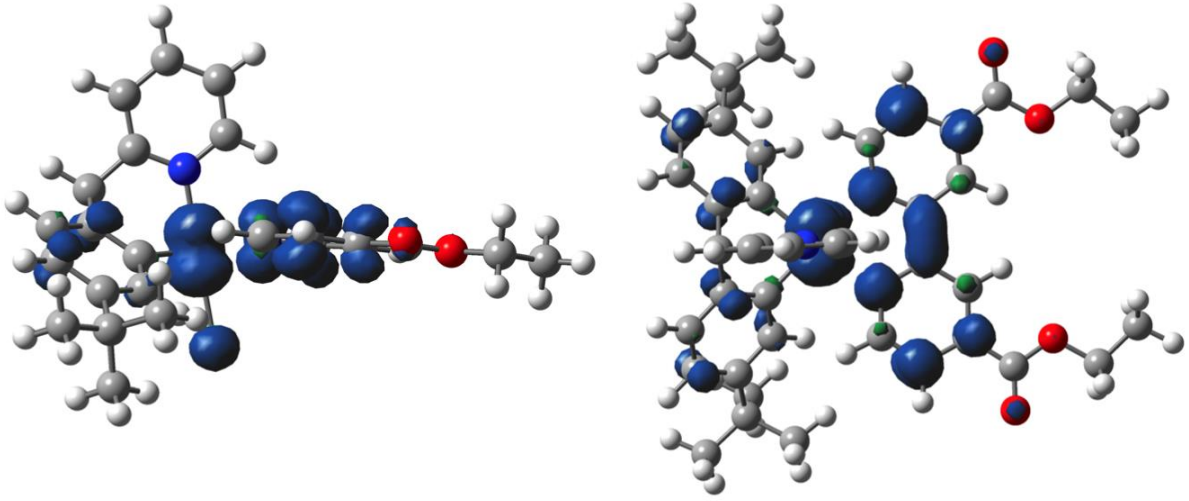


Figure **6.** DFT computed spin density difference plots for the lowest triplet state of **1**. Both side and top views are shown and they have been drawn with a contour threshold of 3×10^{-3} au.

Table **2.** Photophysical properties of complex **1**.

Complex	λ_{abs} / nm (ε / $M^{-1}cm^{-1}$)[a]	λ_{em}[b] / nm	Φ_{PL}[b,c] / %	τ_e[d] / ns
1	237 (34819), 319 (14647), 384 (5105), 434 (5607), 504 (2176), 597 (1925)	731	0.5	36 (73 %) 78 (19 %) 392 (8 %)

[a] Recorded in aerated CH_2Cl_2 at 298 K; [b] Recorded at 298 K in deaerated CH_2Cl_2 solution (λ_{exc} = 420 nm); [c] $[Ru(bpy)_3]PF_6$ in MeCN as the reference (Φ_{PL} = 1.8% in aerated MeCN at 298 K)[50]; [d] λ_{exc} = 378 nm.

Dye-sensitized solar cells (DSSCs)

Sandwich-type solar cells were assembled using **1**-sensitised nanocrystalline TiO_2 as the working electrodes, platinized conducting glass as the counter electrode and iodide/triiodide in acetonitrile as electrolyte. The photovoltaic performances of solar cells based **1** and **N719**, as benchmark sensitizer, are summarized in Table **3**. Figure **7** shows the current–voltage characteristics of the dyes under AM 1.5 simulated sunlight (100 mW cm^{-2}) and in the dark.

Table **3**. Photovoltaic performance of **1**.

DYE	J_{SC}[a] / mA cm^{-2}	V_{OC}[a] / V	**FF**[a]	η[a] / %
1	0.995	0.67	0.74	0.49
N719	8.84	0.81	0.61	4.4

[a] J_{sc} is the short-circuit current density at the $V = 0$ intercept, V_{oc} is the open-circuit voltage at the $J = 0$ intercept, FF is the device fill factor, η is the power conversion efficiency.

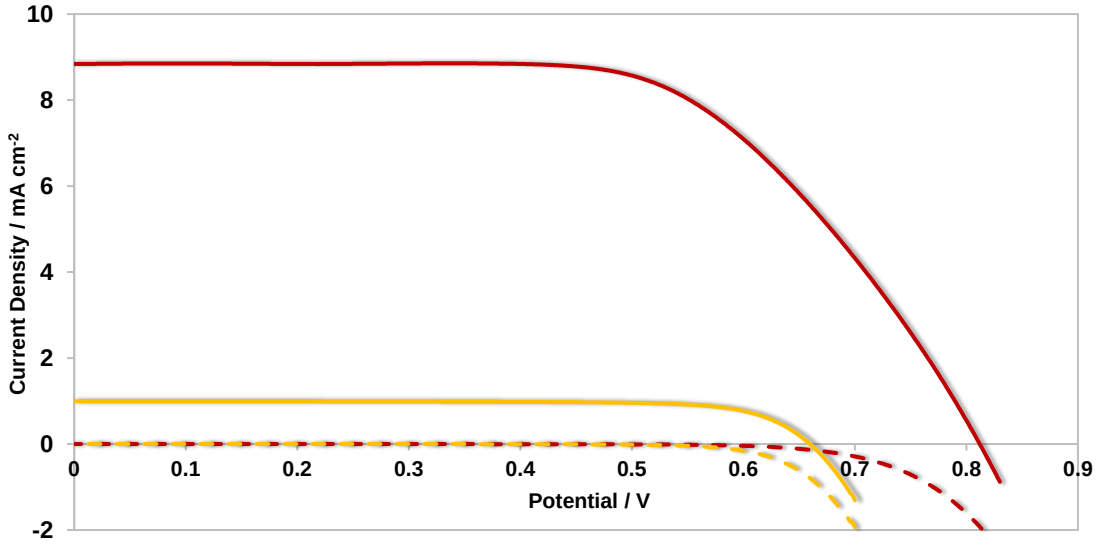


Figure **7**. Current-voltage curves for DSSCs constructed using **1** (orange) and **N719** (red) in the dark (dashed line) and under simulated sunlight (solid line, AM1.5, 100 mW cm^{-2}).

The photovoltaic efficiency ($\eta = 0.49\%$) obtained with **1** is low, but comparable with results for iridium sensitizers reported elsewhere.[10,9,8,7,5] Both charge injection from the excited dye into TiO_2 and regeneration by the electrolyte are thermodynamically favourable. The fill factor for the N719 device (0.61) was slightly lower than typically obtained in optimized devices (0.70-0.75) and the shape of the current-voltage curve is consistent with high series resistance. Procedures used in optimized N719 devices, such as mixed solvents or additives such as chenodeoxycholic acid in the electrolyte or dye bath are likely to improve the performance of the N719 devices, however the conditions chosen were optimal for compound **1**. We therefore attribute the reason for the low efficiency for **1** compared to the benchmark Ru dye to be the weak absorption in the visible region, compared to ruthenium-based photosensitizers such as N719. The absorption spectrum of the TiO_2 electrode after immersion in the dye solution is provided in Figure **S13** and the spectral response of the DSSC is given in Figure **S14**. The low incident photon-to current

conversion efficiency (IPCE < 2%) is consistent with the poor light-harvesting at $\lambda > 500$ nm. While these dyes absorb broadly across the visible spectrum, the low ε ($\varepsilon \sim 2\ 000\ M^{-1}\ cm^{-1}$) compared to ruthenium dyes ($\varepsilon > 10\ 000\ M^{-1}\ cm^{-1}$) is a limitation to their solar cell performance.

Conclusions

In conclusion, a new panchromatically absorbing, NIR luminescent iridium(III) complexes bearing a tripodal tris(six-membered) chelate ligand has been obtained and comprehensively characterized, including by single crystal X-ray diffraction. The absorption spectrum tails off at 700 nm, much further than most neutral iridium complexes while the emission is significantly shifted into the NIR, with a maximum of 731 nm. DSSCs using **1** as the dye achieved only modest efficiency of 0.49%, comparable to other Ir(III) dyes. This was attributed to the modest absorption coefficient, which leads to weak light harvesting in the visible region and low short-circuit current.

Experimental

Dye sensitised solar cell fabrication

FTO glass substrates (TEC 15) were cleaned by ultrasonication in dilute soapy water, 0.1 M HCl in ethanol, and ethanol for 15 minutes each. A $TiCl_4$ blocking layer was deposited by immersing the substrates into 20 mM $TiCl_4$ solution and heating to 75 ^oC for 30 min, then washing with water and ethanol. TiO_2 films were prepared by a doctor blading method in which a transparent TiO_2 paste (DSL 18NR-T, Dyesol) was deposited by spreading with a glass rod. Masks to determine thickness and film size (0.20 cm^2) were made using Scotch Magic Tape. The films were dried by heating to 80 ^oC on a hotplate for 5 minutes. In some cases, a TiO_2 scattering layer (Ti Nanoxide R/SP, Solaronix) was applied. The films were sintered (450 ^oC, 30 min, 30 min ramp time) and

upon cooling, another $TiCl_4$ treatment was applied. The films were placed into dye baths containing 0.3 mM of Complex **1** (the best results were obtained when DCM was used as a solvent), and left in the dark for 24 hours before testing. Control dye baths contained 0.3 mM N719 in a 1:1 ratio of t-butanol and acetonitrile.

The Pt catalyst was deposited on the FTO glass by coating with 10 µL cm^{-2} of H_2PtCl_6 solution (5 mM in isopropanol), followed by heating at 400 °C for 15 minutes. Cells were constructed using the as-prepared films sandwiched with a Pt-coated FTO glass counter electrode (TEC 8), using Surlyn as a spacer. An iodine redox electrolyte composed of 0.6 M TBAI, 0.03 M I_2, 0.1 M guanidinium thiocyanate and 0.5 M t-butyl pyridine in MeCN was introduced into cells via a pre-drilled hole in the counter electrode, which were then sealed with another piece of Surlyn and a glass coverslip. The cells were masked with an aperture 1mm wider than the active area. The current–voltage characteristics of the DSSCs were recorded under AM 1.5 simulated sunlight (100 mW cm^{-2}) and in the dark. The best of a catch of four cells is reported, with results for each set agreeing within 10% for Jsc and 2% for Voc.

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Supporting information. X-ray crystallographic structure of **1** in CIF format (CCDC: 1583853). NMR and MS spectra for **1**, Supplementary crystallographic data, supplementary electrochemical and photophysical data. Description of the DFT/TD-DFT protocol. Experimental details for the DSSC assembly and testing. Plots of the absorption spectra of **1**-sensitized TiO_2 and IPCE spectrum of the DSSC.

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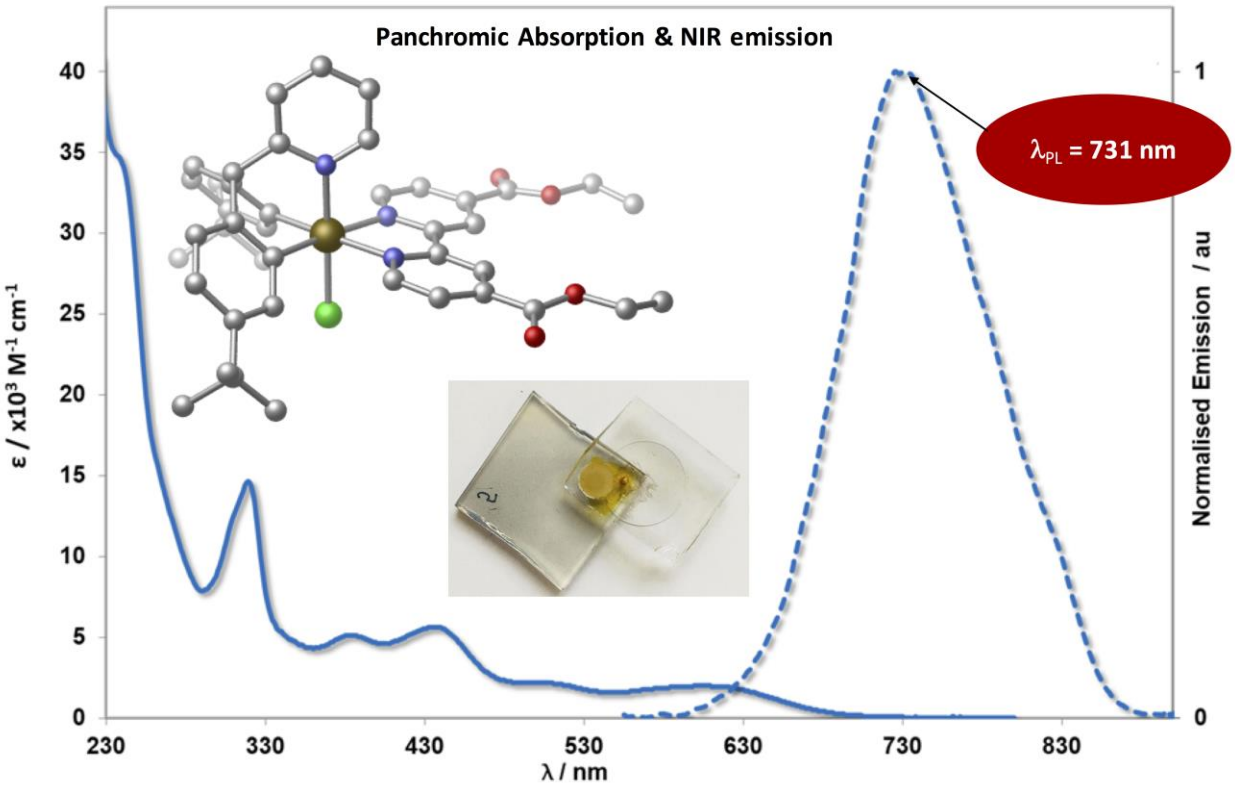
Panchromic Absorption & NIR emission
λPL = 731 nm
ε / x10³ M⁻¹ cm⁻¹
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1
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830